

Purification of caprolactam

5 The present invention relates to a process for removing high boilers from crude caprolactam which comprises high boilers, caprolactam and in some cases low boilers, and which has been obtained by

- a) reacting 6-aminocapronitrile with water to give a reaction mixture
- 10 b) removing ammonia and unconverted water from the reaction mixture to obtain crude caprolactam,

which comprises
- 15 c) feeding the crude caprolactam to a distillation apparatus to obtain a first substream via the top as a product and a second substream via the bottom,
by setting the pressure in the distillation in such a way that the bottom temperature does not go below 170°C, and
adjusting the second substream in such a way that the caprolactam content of the second
20 substream is not less than 10% by weight, based on the entire second substream.

Processes for preparing caprolactam are common knowledge.

25 It is likewise common knowledge, for example from Ullmann's Encyclopedia of Industrial Chemistry, 5th Ed., Vol. A5, VCH Verlagsgesellschaft mbH, Weinheim (Germany), 1986, page 46-48, or Kirk-Othmer, Encyclopedia of Chemical Technology, 4th Ed., Vol. 4, John Wiley & Sons, New York, 1992, page 836, that caprolactam which is used for the preparation of polymers has to have a purity of from 99.9 to 99.94%, and the main impurity is typically water in an amount of from 0.04 to 0.1%. Other impurities may be present only in the region of a
30 maximum of a few ppm.

For instance, caprolactam can be prepared by Beckmann rearrangement of cyclohexanone oxime with sulfuric acid or oleum. After neutralization of the mixture obtained in this way with ammonia, the caprolactam can be obtained from the ammonium sulfate formed as a by-product
35 by extraction with an organic solvent.

Depending on the processes for preparing the reactants used to synthesize cyclohexanone oxime, such as cyclohexanone and hydroxylammonium sulfate, the oximation and rearrangement conditions, the crude caprolactam which has been obtained by Beckmann rearrangement comprises impurities which differ in their type and amount. Typical impurities of

crude caprolactam which has been obtained by Beckmann rearrangement are C-methylcaprolactams, 6-methylvalerolactam and n-pentylacetamide.

5 For the purification of the crude caprolactam obtained in the Beckmann rearrangement, various processes have been described.

According to DE-A-1253716, the crude caprolactam can be purified by hydrogenation in suspension in the presence of a catalyst and with the addition of an acid.

10 According to DE-A-1253716, the crude caprolactam can be purified by hydrogenation in suspension in the presence of a catalyst and with the addition of a base.

DD-A-75083 describes a process for purifying crude caprolactam by initially distilling crude caprolactam and subsequently, dissolved in an organic solvent, hydrogenating in the presence
15 of a catalyst and then treating with an ion exchanger.

According to EP-A-411455, the characteristic important quality features for caprolactam can be maintained by continuously hydrogenating the crude caprolactam in a liquid phase process.

20 Crude caprolactam which is obtained by hydroformylating 3-pentenoic acid and/or its esters to 5-formylvaleric acid/esters as the main products and 4- and 3-formylvaleric acid/esters as the by-products, extractive (WO 97/02228) or distillative (WO 97/06126) removal of these branched formylvaleric acid/esters, aminating hydrogenation of 5-formylvaleric acid/esters to 6-aminocaproic acid/esters and/or 6-aminocapronamide and cyclization of 6-aminocaproic
25 acid/esters or 6-aminocapronamide contains other typical impurities.

For example, the crystallization of crude caprolactam with the addition of 10% by weight of water obtained starting from 5-formylvaleric esters is disclosed by WO 99/48867, Example 1, and starting from mixtures of 6-aminocaproic acid, 6-aminocapronamide and corresponding
30 oligomers by WO 98/37063, Example 9. In this crude caprolactam from which high and low boilers have not been removed before crystallization, 6345 ppm of N-methylcaprolactam, 100 ppm of 5-methylvalerolactam, 78 ppm of valeramide and other impurities were present. The crude caprolactam/water melt impurities were present. The crude caprolactam/water melt was homogenized at 50°C and then cooled to 30°C. The precipitated crystals were filtered off and
35 washed 2 to 3 times with aqueous caprolactam. 5-Methylvalerolactam and valeramide were depleted to 1 ppm, and N-methylcaprolactam to 51 ppm. 73.6 g of crude lactam provided 33.7 g of pure lactam (caprolactam yield: 45.8%). The specification for the volatile bases (VB) was only attained by a second crystallization. When high and low boilers were removed from the crude

caprolactam before the crystallization according to WO 99/48867, Example 3, the caprolactam yield after crystallization was 52%.

5 WO 99/65873 further discloses the selective adsorption of caprolactam from mixtures with 4-ethyl-2-pyrrolidone, 5-methyl-2-piperidone, 3-ethyl-2-pyrrolidone and 3-methyl-2-piperidone or octahydrophenazine on adsorbents such as activated carbon, molecular sieves or zeolites to obtain pure caprolactam after desorption. This caprolactam removal may be followed by a melt crystallization or a crystallization from a solvent.

10 It is also known that crude caprolactam can be purified by crystallization, starting from 6-aminocapronitrile according to WO 98/37063, claim 8, and initially hydrolyzing with water to give 6-aminocaproic acid. Water and ammonia formed by hydrolysis are then removed, the 6-aminocaproic acid formed is cyclized and the resulting crude caprolactam is crystallized according to WO 99/48867.

15 Caprolactam can also be obtained by reacting 6-aminocapronitrile (ACN) with water in the liquid phase in the presence or absence of a catalyst with the release of ammonia.

20 In addition to caprolactam, water, ammonia and in some cases further liquid diluent, the mixture obtained in this reaction comprises impurities having a boiling point above that of caprolactam (high boilers) and those having a boiling point below that of caprolactam (low boilers).

25 The example of US-A-496,941 discloses that crude caprolactam is obtained in a purity of 99.5% after the removal of water, solvent, ammonia, low boilers and high boilers from a mixture obtained in the reaction of ACN with water and solvent.

30 For crude caprolactam which has been obtained from ACN in the liquid phase, other purification processes are described, since the impurities of such crude caprolactam differ significantly from those of crude caprolactam which has been obtained by other processes, as described in US-A-5,496,941.

According to US-A-5,496,941, ACN is converted in a first step in the liquid phase to caprolactam, low boilers, water, ammonia and optionally further solvents are removed simultaneously, high boilers are removed to obtain crude caprolactam in a purity of 99.5%, this
35 crude caprolactam is hydrogenated in the presence of a catalyst, the resulting product is treated with an acidic ion exchanger or sulfuric acid and the resulting product is distilled in the presence of a base.

WO 96/20923 discloses a process for purifying crude caprolactam which stems from liquid phase cyclization of 6-aminocapronitrile with water in the presence of a solvent and of heterogeneous catalysts. In this process, crude caprolactam is initially hydrogenated, then treated with acidic agents and finally distilled in the presence of alkali.

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A disadvantage of these two purification processes is that three separate reaction steps are required for the preparation of pure caprolactam.

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DE 100 21 199 A1 and DE 100 21 192 disclose the purification by crystallization of caprolactam obtained by liquid or gas phase cyclization after removal of ammonia and water.

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The processes mentioned for purifying crude caprolactam which has been obtained from ACN have the disadvantage that they are technically complicated and energy-intensive, in particular as a consequence of the numerous separating steps.

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It is an object of the present invention to provide a process which enables the preparation of caprolactam which has been obtained starting from ACN, in high purity and in a technically simple and energy-saving manner.

We have found that this object is achieved by the process defined at the outset.

In the process according to the invention, crude caprolactam is used which has been obtained by reacting 6-aminocapronitrile with water according to steps a) and b).

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In step a), a mixture (I) comprising 6-aminocapronitrile, water and optionally liquid diluent is converted to a mixture (II) comprising caprolactam, ammonia, water, optionally liquid diluent, high boilers and optionally low boilers, preferably in the presence of a solid which catalytically supports the conversion.

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The ACN required for step a) can, as generally known from Ullmann's Encyclopedia of Industrial Chemistry, 5th Ed., Vol. A5, VCH Verlagsgesellschaft mbH, Weinheim (Germany), 1986, page 46, Fig. 8, be obtained from adiponitrile.

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Particular useful processes are the partial catalyzed hydrogenation of adiponitrile in the presence of ammonia as a solvent and, for example as a suspension catalyst, rhodium on magnesium oxide (US-A-4,601,859), Raney nickel (US-A-2,762,835, WO 92/21650), nickel on aluminum oxide (US-A-2,208,598), or, as a fixed bed catalyst, Cu-Co-Zn spinel (DE-B-954416, US-A-2,257,814) or iron (DE-A-42 35 466) or a process according to US-A-2,245,129, US-A-2,301,964, EP-A-150295, FR-A-2 029 540 or a process described in US-A-5,496,941.

The adiponitrile required for this conversion is prepared industrially, for example, by double hydrocyanation of butadiene in the presence of nickel-containing catalysts and is commercially available, for example, from Aldrich-Chemie Gesellschaft mbH & Co. KG, Steinheim, Germany.

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According to US-A-4,628,085, mixture (I) can be converted to mixture (II) in the gas phase over silica gel at 300°C.

Equally, this conversion can be carried out according to US-A-4,625,023 in the gas phase over a silica gel or copper/chromium/barium-titanium oxide catalyst.

- 10 According to FR-A-2029540, the reaction can be carried out in the presence of catalysts which are metallic Zn or Cu powder or oxides, hydroxides, halides, cyanides of rubidium, lead, mercury or of the elements having an atomic number of from 21 to 30 or from 39 to 48. The catalysts described are used as suspension catalysts in batchwise stirred autoclaves.

- 15 Mixture (I) can also be converted to mixture (II), for example, according to EP-A-659 741, WO 96/22974, DE 19632006, WO 99/47500 or WO 99/28296.

- The conversion can preferably be carried out in the gas phase at temperatures of generally from 200 to 550°C, preferably from 250 to 400°C; the pressure is generally in the range from 0.01 to 20 bar, preferably at atmospheric pressure, although care has to be taken that the reaction mixture is predominantly gaseous under the conditions employed.

- The catalyst hourly space velocities are typically from 0.05 to 2 kg, preferably from 0.1 to 1.5 kg, in particular from 0.2 to 1 kg, of 6-aminocapronitrile per liter of catalyst volume per hour.

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The conversion may be carried out batchwise, but preferably continuously.

- Useful reactors are advantageously those as generally known for gas phase reactions over moving or stationary solid catalysts. Preference is given to using a fluidized bed reactor, 30 preferably fixed bed reactor, such as a tray reactor, in particular a tubular reactor. Combinations of such reactors are also possible.

Generally from 1 to 50 mol, preferably from 1 to 10 mol, of water are used per mole of ACN.

- 35 The mixture (I) may also comprise further organic compounds which are in gaseous form under the reaction conditions, such as alcohols, amines or aromatic or aliphatic hydrocarbons.

Useful catalytically active compounds of the catalysts are, for example, silicon dioxide as pyrogenic silicon dioxide, as silica gel, kieselguhr, quartz or mixtures thereof, copper chromite, preferably aluminum oxide, titanium oxide, preferably titanium dioxide, lanthanum phosphates, lanthanum oxides, and also mixtures of such compounds.

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Aluminum oxide is suitable in all modifications which can be obtained by heating the precursor compounds aluminum hydroxide (gibbsite, boehmite, pseudoboehmite, bayerite and diaspor) at different temperatures. These include in particular gamma- and alpha-aluminum oxide and their mixtures.

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Titanium dioxide is amorphous and suitable in all its modifications, preferably anatase and rutile, and also mixtures of such modifications.

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Lanthanum phosphates are suitable in their various modifications, stoichiometric ratios between lanthanum and phosphate units and degrees of condensation of the phosphate units (monophosphate, oligophosphates such as diphosphates or triphosphates, polyphosphates), individually or in a mixture.

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These compounds can be used in the form of powders, meal, spall, extrudates or pressed to tablets. The form of the compounds is generally determined by the requirements of the particular reaction method, and powder or meal are advantageously used in the fluidized bed method. In the fixed bed method, tablets or extrudates having diameters of between 1 mm and 6 mm are customarily used.

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The compounds can be used in pure form (content of the particular compounds > 80% by weight), as a mixture of the abovementioned compounds, in which case the sum of the abovementioned compounds should be > 80% by weight, or as a supported catalyst, in which case the abovementioned compounds can be applied to a mechanically and chemically stable support, usually having a high surface area.

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The pure compounds may have been prepared by precipitating from aqueous solutions, for example titanium dioxide by the sulfate process or by other processes such as pyrogenic preparation of fine aluminum oxide, titanium dioxide or zirconium dioxide powders, which are commercially available.

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Several methods are available for preparing the mixtures of various compounds. The compounds or their precursor compounds which can be converted to the oxides by calcining can be prepared, for example, by coprecipitation from solution. This generally results in very good

distribution of the two compounds used being obtained. The compound or precursor mixtures can also be precipitated by precipitating one compound or precursor in the presence of finely divided particles of the second compound or precursor present as a suspension. A further method consists in mechanical mixing of the compound or precursor powder, and this mixture may find use as a starting material for producing extrudates or tablets.

All methods described in the literature are in principle suitable for preparing supported catalysts. For instance, the compounds can be applied to the support in the form of their sols by simple impregnation. Drying and calcining customarily remove the volatile constituents of the sol from the catalyst. Such sols are commercially available for titanium dioxide and aluminum oxide.

A further possibility for applying layers of the catalytically active compounds is to hydrolyze or pyrolyze organic or inorganic compounds. For instance, a ceramic support can be covered with a thin layer of titanium dioxide by hydrolysis of titanium isopropoxide or other titanium alkoxides. Further suitable compounds include TiCl_4 and aluminum nitrate. Suitable supports are powders, extrudates or tablets of the compounds mentioned themselves or other stable compounds such as steatite or silicon carbide. The supports used may be macroporous to improve the mass transfer.

The reaction can be carried out in the presence of a gas inert with regard to the conversion of mixture (I) to mixture (II), preferably argon, in particular nitrogen. The volume ratio of inert gas to the ACN which is gaseous under the reaction conditions may advantageously be up to 100.

Particular preference is given to step 1) being a process as described in US-A-5,646,277 or US-A-5,739,324.

In these processes, the reaction is carried out in the liquid phase at temperatures of generally from 140 to 320°C, preferably from 160 to 280°C; the pressure is generally in the range from 1 to 250 bar, preferably from 5 to 150 bar, although care has to be taken that the reaction mixture is predominantly liquid under the conditions employed. The residence times are generally in the range from 1 to 120 min, preferably from 1 to 90 min and in particular from 1 to 60 min. In some cases, residence times of from 1 to 10 min have proven entirely sufficient.

The conversion can be carried out batchwise, but preferably continuously. Useful reactors include a stirred tank, autoclave, preferably a fixed bed tubular reactor. Combinations of such reactors are also possible.

Generally at least 0.1 mol, preferably from 0.5 to 100 mol and in particular from 1 to 20 mol, of water are used per mole of ACN.

Advantageously, the ACN is used in the form of a from 1 to 50% by weight, in particular from 5 to 50% by weight, more preferably from 5 to 30% by weight, solution in water, in which case the solvent is then at the same time a reaction partner, or in mixtures comprising water and a liquid diluent. Examples of useful diluents include alkanols, such as methanol, ethanol, n- and i-propanol, n-, i- and t-butanol, and polyols such as diethylene glycol and tetraethylene glycol, hydrocarbons such as petroleum ether, benzene, toluene, xylene, lactams such as pyrrolidone or caprolactam or alkyl-substituted lactams such as N-methylpyrrolidone, N-methylcaprolactam or N-ethylcaprolactam and also carboxylic esters, preferably of carboxylic acids having from 1 to 8 carbon atoms. Ammonia may also be present in the reaction. It will be appreciated that mixtures of organic liquid diluents may also find use. Mixtures of water and alkanols in a water/alkanol weight ratio of 1-75/25-99, preferably 1-50/50-99 have in some cases proven particularly advantageous.

It is in principle also possible to employ ACN as a reactant and at the same time to employ solvents.

In a particularly preferred embodiment, useful liquid diluents are those which have a miscibility gap with water under certain pressure, temperature and concentration conditions. In this context, a miscibility gap is the separation of the mixture into two liquid phases, one of the two phases having a higher proportion by weight of water, based on the sum of water and liquid diluent, than the other phase. Particularly suitable liquid diluents therefor are hydrocarbons, such as benzene, toluene or xylene, in particular toluene.

The reaction of step a) may be carried out in the miscibility gap, i.e. in the presence of two liquid phases, or preferably outside the miscibility gap, i.e. in the presence of one liquid phase.

Useful heterogeneous catalysts are, for example: acids, basic or amphoteric oxides of the elements of the second, third or fourth main group of the Periodic Table, such as calcium oxide, magnesium oxide, boron oxide, aluminum oxide, tin oxide or silicon dioxide as pyrogenic silica, as silica gel, kieselguhr, quartz or mixtures thereof, and also oxides of metals of the second to sixth transition group of the Periodic Table, such as amorphous titanium oxide, as anatase or rutile, zirconium oxide, zinc oxide, manganese oxide or mixtures thereof. It is likewise possible to use oxides of lanthanides and actinides, such as cerium oxide, thorium oxide, praseodymium oxide, samarium oxide, rare earth mixed oxides, or mixtures thereof with the aforementioned

oxides. Further catalysts may, for example, be:

vanadium oxide, niobium oxide, iron oxide, chromium oxide, molybdenum oxide, tungsten oxide or mixtures thereof. Mixtures of the oxides mentioned with each other are likewise possible.

- 5 Some sulfides, selenides and tellurides, such as zinc telluride, zinc selenide, molybdenum sulfide, tungsten sulfide, sulfides of nickel, zinc and chromium, can also be used.

The aforementioned compounds may be doped with compounds of the 1st and 7th main group of the Periodic Table or contain them.

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Further suitable catalysts include zeolites, phosphates and heteropolyacids, and also acidic and alkaline ion exchangers, for example Nafion.

- 15 These catalysts may optionally each comprise up to 50% by weight of copper, tin, zinc, manganese, iron, cobalt, nickel, ruthenium, palladium, platinum, silver or rhodium.

- 20 Particularly preferred catalysts which have very high yields, conversions, selectivities and on-stream times under the above-described reaction conditions are heterogeneous catalysts based on titanium oxide, zirconium oxide, cerium oxide and aluminum oxide. These may be used in the form of powders, meal, spall, extrudates or pressed to tablets. The form of the oxides is generally determined by the requirements of the particular reaction method, and powder or meal are used in suspension. In the fixed bed method, it is customary to use tablets or extrudates having diameters between 1 mm and 10 mm.

- 25 Aluminum oxide is suitable in all modifications which can be obtained by heating the precursor compounds aluminum hydroxide (gibbsite, boehmite, pseudoboehmite, bayerite and diasporite) at different temperatures. These include in particular gamma- and alpha-aluminum oxide and their mixtures.

- 30 The oxides can be used in pure form (content of the particular oxide > 80% by weight), as a mixture of the abovementioned oxides, in which case the sum of the abovementioned oxides should be > 80% by weight, or as a supported catalyst, in which case the abovementioned oxides can be applied to a mechanically and chemically stable support, usually having a high surface area.

35

The pure oxides may have been prepared by precipitating from aqueous solutions, for example titanium dioxide by the sulfate process or by other processes such as pyrogenic preparation of fine aluminum oxide, titanium dioxide or zirconium dioxide powders, which are commercially

available.

Several methods are available for preparing the mixtures of various oxides. The oxides or their precursor compounds which can be converted to the oxides by calcining can be prepared, for example, by coprecipitation from solution. This generally results in very good distribution of the two oxides used being obtained. The oxide or precursor mixtures can also be precipitated by precipitating one oxide or precursor in the presence of finely divided particles of the second oxide or precursor present as a suspension. A further method consists in mechanical mixing of the oxide or precursor powder, and this mixture may find use as a starting material for producing extrudates or tablets.

All methods described in the literature are in principle suitable for preparing supported catalysts. For instance, the oxides can be applied to the support in the form of their sols by simple impregnation. Drying and calcining customarily remove the volatile constituents of the sol from the catalyst. Such sols are commercially available for titanium dioxide, aluminum oxide and zirconium dioxide.

A further possibility for applying layers of the active oxides is to hydrolyze or pyrolyze organic or inorganic oxides. For instance, a ceramic support can be covered with a thin layer of titanium dioxide by hydrolysis of titanium isopropoxide or other titanium alkoxides. Further suitable oxides include TiCl_4 , zirconyl chloride, aluminum nitrate and cerium nitrate. Suitable supports are powders, extrudates or tablets of the oxides mentioned themselves or other stable oxides such as silicon dioxide. The supports used may be macroporous to improve the mass transfer.

In step b), ammonia and unconverted water are removed from mixture (II) to obtain a mixture (III) comprising caprolactam, any liquid diluent, high boilers and low boilers.

The ammonia can in principle be removed from mixture (II) by processes known per se for separation of materials, such as extraction or preferably distillation, or a combination of such processes.

The distillation may advantageously be carried out at bottom temperatures of from 60 to 220°C, in particular from 100 to 220°C. It is customary to set a pressure, measured at the top of the distillation apparatus, of from 2 to 30 bar absolute.

Useful apparatus for the distillation is apparatus customary for this purpose, as described, for example, in Kirk-Othmer, Encyclopedia of Chemical Technology, 3rd Ed., Vol. 7, John Wiley &

Sons, New York, 1979, page 870-881, such as sieve tray columns, bubble-cap tray columns, and columns having structured packing or random packing.

5 The distillation can be carried out in a plurality of, such as 2 or 3 columns, advantageously a single column.

In step b), water, fully or partly, and any liquid diluent are removed from mixture (III) to obtain crude caprolactam (IV) comprising caprolactam, high boilers and optionally low boilers.

10 When a liquid diluent has been used in step a), water and liquid diluent can be removed in step b) simultaneously, or the water can be removed before or after the liquid diluent.

The water and any liquid diluent from mixture (III) can in principle be removed by processes known per se for separation of materials, such as extraction, crystallization or preferably
15 distillation, or a combination of such processes.

The distillation may advantageously be carried out at bottom temperatures of from 50 to 250°C, in particular from 100 to 230°C.

20 Useful apparatus for the distillation is apparatus customary for this purpose, as described, for example, in Kirk-Othmer, Encyclopedia of Chemical Technology, 3rd Ed., Vol. 7, John Wiley & Sons, New York, 1979, page 870-881, such as sieve tray columns, bubble-cap tray columns, and columns having structured packing or random packing.

25 Particular preference is given to heat-integrated, multistage removal of the water and of any liquid diluent.

When, in a particularly preferred embodiment, step a) is carried out in the presence of a liquid diluent which has a miscibility gap with water, the water can advantageously be removed by
30 converting the system, especially when carrying out step a) in a single liquid phase, into this miscibility gap to form two liquid phases, of which one phase has a higher proportion by weight of water, based on the sum of water and liquid diluent, than the other phase, and then removing the phase with the higher proportion by weight of water from the other phase which typically contains the predominant amount of crude caprolactam.

35 The liquid diluent can be removed from the phase comprising the predominant amount of crude caprolactam by processes known per se, for example as already described above.

Before feeding the crude caprolactam (IV) into step c), in a preferred embodiment, the low boilers are removed from the crude caprolactam (IV) between steps b) and c).

5 The low boilers can in principle be removed by processes known per se for separation of materials, such as extraction, crystallization or preferably distillation, or a combination of such processes.

10 The distillation may advantageously be carried out at bottom temperatures of from 50 to 250°C, in particular from 100 to 230°C. It is customary to set a pressure, measured at the top of the distillation apparatus, of from 1 to 500 mbar absolute, preferably from 5 to 100 mbar absolute.

15 Useful apparatus for the distillation is apparatus customary for this purpose, as described, for example, in Kirk-Othmer, Encyclopedia of Chemical Technology, 3rd Ed., Vol. 7, John Wiley & Sons, New York, 1979, page 870-881, such as sieve tray columns, bubble-cap tray columns, and columns having structured packing or random packing.

The distillation for removing the low boilers can be carried out in a plurality of, such as 2 or 3 columns, advantageously a single column.

20 The distillation for removing the high boilers can be carried out in a plurality of, such as 2 or 3 columns, advantageously a single column.

The low boilers in this removal are in particular 6-aminocapronitrile.

25 In a further embodiment, the low boilers are removed from the product stream after step c).

The low boilers can in principle be removed by processes known per se for separation of materials, such as extraction, crystallization or preferably distillation, or a combination of such processes.

30 The distillation may advantageously be carried out at bottom temperatures of from 50 to 250°C, in particular from 100 to 230°C. It is customary to set a pressure, measured at the top of the distillation apparatus, of from 1 to 500 mbar absolute, preferably from 5 to 100 mbar absolute.

35 Useful apparatus for the distillation is apparatus customary for this purpose, as described, for example, in Kirk-Othmer, Encyclopedia of Chemical Technology, 3rd Ed., Vol. 7, John Wiley & Sons, New York, 1979, page 870-881, such as sieve tray columns, bubble-cap tray columns, and columns having structured packing or random packing.

The distillation for removing the low boilers can be carried out in a plurality of, such as 2 or 3 columns, advantageously a single column.

5 The distillation for removing the high boilers can be carried out in a plurality of, such as 2 or 3, columns, advantageously a single column.

The low boilers in this removal are in particular 6-aminocapronitrile.

10 According to the invention, the crude caprolactam is fed to a distillation apparatus C1.

In this distillation apparatus, a first substream is obtained via the top. This substream comprises substantially purified caprolactam as the product.

15 In this distillation apparatus, a second substream is also obtained via the bottom. This substream comprises caprolactam and high boilers.

20 Useful distillation apparatus is apparatus customary for this purpose, as described, for example, in: Kirk-Othmer, Encyclopedia of Chemical Technology, 3rd Ed., Vol. 7, John Wiley & Sons, New York, 1979, page 870-881, such as sieve tray columns, bubble-cap tray columns, and columns having structured packing or random packing.

The distillation may be carried out in a plurality of, such as 2 or 3, columns, advantageously a single column.

25 When a column is used as the first distillation apparatus, this column may advantageously have from 1 to 30 theoretical plates, in particular from 5 to 20 theoretical plates.

30 In a preferred embodiment, crude caprolactam (IV) can be introduced to the top of the distillation apparatus C1.

According to the invention, the pressure in the distillation apparatus C1 is such that the bottom temperature does not go below 170°C, preferably 185°C.

35 An advantageous bottom temperature is at most 300°C, in particular at most 250°C.

Moreover, the second substream is adjusted in such a way that the caprolactam content of the second substream is not less than 10% by weight, preferably not less than 15% by weight, in particular not less than 20% by weight, based on the entire second substream.

- 5 The parameters for the operation of the selected distillation apparatus which are optimum for the achievement of the provisions according to the invention can be easily determined by a few simple preliminary experiments.

In a preferred embodiment, the second substream can be partly or fully recycled in step a).

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It was surprising that the invention succeeds in distillatively converting crude caprolactam prepared from ACN in a technically simple and economic manner to caprolactam from which polycaprolactam can be prepared.

- 15 Examples
 Example 1

105 kg/h of crude caprolactam which had been obtained by reacting 6-aminocapronitrile with water were fed to a distillation column having 5 theoretical plates.

- 20 85 kg/h of caprolactam were removed via the top.

20 kg/h of caprolactam having a high boiler content of 25% by weight, based on the entire bottom stream, were removed via the bottom.

The bottom temperature was 190°C.

- 25 The distillation column could be operated uninterrupted under these conditions for more than one month.

The caprolactam obtained via the top was colorless and could be polymerized without any problem.

Comparative example 1

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The procedure of example 1 was repeated with the variation that the bottom takeoff amount was reduced in such a way that the high boiler content was 80% by weight, based on the entire bottom stream.

The distillation apparatus could initially be operated.

- 35 However, after two days, solids had formed in the bottom which blocked the lines and prevented the further operation of the distillation column.

Comparative example 2

The procedure of example 1 was repeated, with the variation that the bottom temperature was only 150°C.

- 5 The distillation apparatus could initially be operated.
However, after 12 hours, solids had formed in the bottom which blocked the lines and prevented the further operation of the distillation column.